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Genetic test in MEN1 syndrome: An evolving story

Francesca Marini, Francesca Giusti, Maria Luisa Brandi

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Abstract

Multiple endocrine neoplasia type 1 (MEN1) is an autosomal dominant inherited tumour syndrome expressing various endocrine and non-endocrine lesions and tumours. Since the identification of the causative gene, the oncosuppressor gene *MEN1*, in 1997, genetic testing has revealed an important approach for the early and differential diagnosis of the disease. 7 The finding of a *MEN1* mutation in a patient has important clinical implications for relatives since it allows very early disease diagnosis and identification of carriers, even

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